L Nun	mber	Hits	Search Text	DB	Time stamp
2		965	replicon and cap	USPAT;	2004/10/28 15:10
				US-PGPUB;	
				EPO; JPO;	
				DERWENT	
3		_ 25	replicon same cap	USPAT;	2004/10/28 15:02
				US-PGPUB;	
ĺ				EPO; JPO;	
4			and the second second (UDD COM)	DERWENT	0004/10/00 15
4		205	replicon and cap and (VEE or SFV or Sindbis or sin)	USPAT;	2004/10/28 15:04
			Sindbis of sin	US-PGPUB; EPO; JPO;	
				DERWENT	
5 .		23	replicon and (helper same cap) and (VEE or	USPAT;	2004/10/28 15:05
			SFV or Sindbis or sin)	US-PGPUB;	2001/10/20 13:03
			,	EPO; JPO;	
				DERWENT	
6		58	replicon and uncapped	USPAT;	2004/10/28 15:11
				US-PGPUB;	İ
ŀ				EPO; JPO;	
				DERWENT	
7		18	replicon and uncapped and (VEE or sin or	USPAT;	2004/10/28 15:11
			sfv or sindbis)	US-PGPUB;	
			*	EPO; JPO;]
_		` 5502	replicon	DERWENT	2004/04/00 33 55
_		3502	replicon	USPAT;	2004/04/29 11:56
			,	US-PGPUB; EPO; JPO;	
				DERWENT	
_		133	(alphavir\$ or (alpha adj vir\$)) same	USPAT;	2004/04/29 11:56
			replicon	US-PGPUB;	2001/01/23 11:30
				EPO; JPO;	
			· .	DERWENT	
_		22638	helper	USPAT;	2004/04/29 11:57
				US-PGPUB;	
				EPO; JPO;	
				DERWENT	
-		42		USPAT;	2004/04/29 13:39
			replicon) same helper	US-PGPUB;	
				EPO; JPO;	
_		4	alphaviral adj replicon	DERWENT	2004/04/29 12:50
		-	aiphavitai adj lepticon	USPAT; US-PGPUB;	2004/04/29 12:50
				EPO; JPO;	
			·	DERWENT	
-		562	rayner.in.	USPAT;	2004/04/29 13:41
				US-PGPUB;	
			<u>'</u>	EPO; JPO;	
				DERWENT	
_		10	kamrud.in.	USPAT;	2004/04/29 16:54
				US-PGPUB;	
				EPO; JPO;	
_	.	23675	ionia adi strongth	DERWENT	2004/04/00 15 55
		23013	ionic adj strength	USPAT;	2004/04/29 16:55
				US-PGPUB; EPO; JPO;	~}
			· ·	DERWENT	`
_		205	(ionic adj strength) same virus	USPAT;	2004/04/29 16:56
				US-PGPUB;	, 51, 25 10, 50
				EPO; JPO;	
				DERWENT	
-		106	((ionic adj strength) same virus).ab,bsum.	USPAT;	2004/04/29 17:49
			•	US-PGPUB;	
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		7.00		DERWENT	
_		7882	johnston.in.	USPAT;	2004/04/29 17:49
				US-PGPUB;	
				EPO; JPO;	
				DERWENT	L

-	4	michael adj3 johnston.in.	USPAT;	2004/04/29 17:50
			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	6	johnston-michael-d.in.	USPAT;	2004/04/29 17:51
`		'	US-PGPUB;	
			EPO; JPO;	
			DERWENT	1
-	5	johnston-michael-denis.in.	USPAT;	2004/04/29 17:51
			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	0	("5204257A").PN.	USPAT	2004/04/30 08:56
-	1	("5204257").PN.	USPAT	2004/04/30 09:00
-	1	("6267967").PN.	USPAT	2004/04/30 11:02
-	5	("3985615" "5024836" "5360736"	USPAT	2004/04/30 09:00
		"5447859" "5607852").PN.		
-	1	("5185440").PN.	USPAT	2004/04/30 11:34
-	1	("5643576").PN.	USPAT	2004/04/30 12:43
-	146	(424/218.1).CCLS.	USPAT	2004/04/30 12:45
-	412	alphavirus	USPAT	2004/04/30 12:46
-	25	((424/218.1).CCLS.) and alphavirus	USPAT	2004/04/30 12:47
-	442	(435/239).CCLS.	USPAT	2004/04/30 12:48
-	5	alphavirus and ((435/239).CCLS.)	USPAT	2004/04/30 13:49
-	3338	replicon	USPAT	2004/04/30 13:50
-	20679	electropo\$	USPAT	2004/04/30 13:50
-	1766	replicon and electropo\$	USPAT	2004/04/30 13:50
-	412	alphavirus	USPAT	2004/04/30 13:51
-	65	(replicon and electropo\$) and alphavirus	USPAT	2004/04/30 13:55
_	3338		USPAT	2004/04/30 13:56
_	65	((replicon and electropo\$) and alphavirus)	USPAT	2004/04/30 14:10
	1	and replicon		
-	1	5185440.pn.	USPAT	2004/04/30 14:13
_	1	electropo\$ and 5185440.pn.	USPAT	2004/04/30 14:17
_	1	5792462.pn.	USPAT	2004/04/30 14:18
-	677	venezuelan	USPAT	2004/04/30 14:18
-	1	5792462.pn. and venezuelan	USPAT	2004/04/30 14:18

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SINCE FILE

ENTRY

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TOTAL SESSION

0.21

FILE 'MEDLINE' ENTERED AT 12:25:26 ON 28 OCT 2004

FILE LAST UPDATED: 27 OCT 2004 (20041027/UP). FILE COVERS 1950 TO DATE.

On February 29, 2004, the 2004 MeSH terms were loaded. See HELP RLOAD for details.

OLDMEDLINE now back to 1950.

MEDLINE thesauri in the /CN, /CT, and /MN fields incorporate the MeSH 2004 vocabulary. See http://www.nlm.nih.gov/mesh/ and http://www.nlm.nih.gov/pubs/techbull/nd03/nd03_mesh.html for a description of changes.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s replicon and cap

2841 REPLICON

1132 REPLICONS

3303 REPLICON

(REPLICON OR REPLICONS)

15208 CAP

4347 CAPS

17808 CAP

(CAP OR CAPS)

L1 15 REPLICON AND CAP

=> dis ti 1-15

- L1 ANSWER 1 OF 15 MEDLINE on STN
- TI Structural properties of a multifunctional T-shaped RNA domain that mediate efficient tomato bushy stunt virus RNA replication.
- L1 ANSWER 2 OF 15 MEDLINE on STN
- TI In vitro replication of hepatitis E virus (HEV) genomes and of an HEV replicon expressing green fluorescent protein.
- L1 ANSWER 3 OF 15 MEDLINE on STN
- TI Inhibitor RNA blocks the protein translation mediated by hepatitis C virus internal ribosome entry site in vivo.
- L1 ANSWER 4 OF 15 MEDLINE on STN
- TI The regulation of hepatitis C virus (HCV) internal ribosome-entry site-mediated translation by HCV replicons and nonstructural proteins.
- L1 ANSWER 5 OF 15 MEDLINE on STN
- TI Hepatitis C virus subgenomic replicons induce endoplasmic reticulum stress activating an intracellular signaling pathway.
- L1 ANSWER 6 OF 15 MEDLINE on STN
- TI Genetic analysis of a poliovirus/hepatitis C virus chimera: new structure for domain II of the internal ribosomal entry site of hepatitis C virus.
- L1 ANSWER 7 OF 15 MEDLINE on STN
- TI A stem-loop motif formed by the immediate 5' terminus of the bovine viral diarrhea virus genome modulates translation as well as replication of the viral RNA.
- L1 ANSWER 8 OF 15 MEDLINE on STN
- TI Characterization of the initiation sites of both polarity strands of a viroid RNA reveals a motif conserved in sequence and structure.

T.1 ANSWER 9 OF 15 MEDLINE on STN TI Foot-and-mouth disease virus 3C protease induces cleavage of translation initiation factors eIF4A and eIF4G within infected cells. MEDLINE on STN L1ANSWER 10 OF 15 TI Uncoupled expression of p33 and p92 permits amplification of tomato bushy stunt virus RNAs. ANSWER 11 OF 15 MEDLINE on STN L1The -45 region of the Escherichia coli lac promoter: CAP ΤI -dependent and CAP-independent transcription. L1ANSWER 12 OF 15 MEDLINE on STN TI Decoying the cap- mRNA degradation system by a double-stranded RNA virus and poly(A) - mRNA surveillance by a yeast antiviral system. L1ANSWER 13 OF 15 MEDLINE on STN Evidence that the SKI antiviral system of Saccharomyces cerevisiae acts by TI blocking expression of viral mRNA. L1ANSWER 14 OF 15 MEDLINE on STN TI Involvement of cell shape in the replication and segregation of chromosomes in Escherichia coli. ANSWER 15 OF 15 MEDLINE on STN L1TIMapping of the multiple regulatory sites for putP and putA expression in the putC region of Escherichia coli. => s cap helper 15208 CAP 4347 CAPS 17808 CAP (CAP OR CAPS) 27790 HELPER 962 HELPERS 28522 HELPER (HELPER OR HELPERS) L2 0 CAP HELPER (CAP(W) HELPER) => s cap and helper 15208 CAP 4347 CAPS 17808 CAP (CAP OR CAPS) 27790 HELPER 962 HELPERS 28522 HELPER (HELPER OR HELPERS) L366 CAP AND HELPER => s 13 and (VEE or Sindbis or SFV) 366 VEE 1 VEES 367 VEE (VEE OR VEES) 2081 SINDBIS

> 968 SFV 41 SFVS 971 SFV

L4

(SFV OR SFVS)

1 L3 AND (VEE OR SINDBIS OR SFV)

ANSWER 1 OF 1 MEDLINE on STN

TIA cis-acting mutation in the Sindbis virus junction region which affects subgenomic RNA synthesis.

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ANSWER 1 OF 1 MEDLINE on STN

A cis-acting mutation in the Sindbis virus junction region which

affects subgenomic RNA synthesis.

ACCESSION NUMBER: DOCUMENT NUMBER:

90064787 MEDLINE PubMed ID: 2685355

TITLE:

A cis-acting mutation in the Sindbis virus

junction region which affects subgenomic RNA synthesis.

AUTHOR: Grakoui A; Levis R; Raju R; Huang H V; Rice C M

CORPORATE SOURCE: Department of Molecular Microbiology, Washington University

School of Medicine, St. Louis, Missouri 63110-1093.

CONTRACT NUMBER: AI24134 (NIAID)

AI26763 (NIAID)

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Journal of virology, (1989 Dec) 63 (12) 5216-27.

Journal code: 0113724. ISSN: 0022-538X.

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DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

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FILE SEGMENT: Priority Journals

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Last Updated on STN: 19980206 Entered Medline: 19891227

The synthesis of Sindbis virus minus-strand and genomic and AB subgenomic RNAs is believed to require specific cis-acting sequences or structures in the template RNAs and a combination of virus-specific proteins and host components which act in trans. A conserved sequence of about 21 nucleotides in the junction region and encompassing the start site for the subgenomic RNA has been proposed to function as the promoter on the minus-strand template for synthesis of the subgenomic RNA (J.-H. M. Rice, L. Dalgarno, E. G. Strauss, and J. H. Strauss, Proc. Natl. Acad. Sci. USA 79:5235-5239, 1982). We introduced a three-base insertion in this sequence, which also inserts a single amino acid near the COOH terminus of nsP4, in a cDNA clone of Sindbis virus from which infectious RNA transcripts can be generated. phenotype of this mutant, called Toto1100CR4.1, was studied after RNA transfection of chicken embryo fibroblasts or BHK cells. The mutation leads to a drastic reduction in the level of the subgenomic RNA but does not alter the start site of the RNA. Probably as a consequence of depressed structural-protein synthesis, very few progeny virions are released and the mutant makes tiny or indistinct plaques even after prolonged incubation. The cis-acting effect of this mutation was demonstrated by incorporating either a wild-type or mutant junction region into a defective-interfering RNA and examining the relative synthesis of defective-interfering RNA-derived subgenomic RNA in vivo in the presence of wild-type helper virus. These results show that the junction region is recognized by yet unidentified viral trans-acting components for subgenomic RNA synthesis. When the Totol100CR4.1 mutant was passaged in culture, plaque morphology variants readily arose. A total of 24 independent revertants were isolated, and 16 were characterized in detail. All revertants analyzed showed an increase in the level of subgenomic RNA synthesis. Sequence analysis of the junction region showed that all were pseudorevertants, with only two containing potentially compensating changes in the junction region. An assay was developed to identify revertants with second-site changes in trans-acting viral components involved in subgenomic RNA synthesis. At least two such revertants were

identified. Mapping of these and other second-site compensating mutations may provide genetic clues as to which virus-specific protein(s) is responsible for interaction with the conserved junction region to promote subgenomic RNA synthesis.

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Connection closed by remote host END

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